



# The reproductive response of the sea urchins *Paracentrotus lividus* (G.) and *Psammechinus miliaris* (L.) to an hyperproteinated macrophytic diet

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1 **The reproductive response of the sea urchins Paracentrotus**  
 2 **lividus (G.) and Psammechinus miliaris (L.) to an**  
 3 **hyperproteinated macrophytic diet.**

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15

16 **Abstract**

17

18 The sea urchins Paracentrotus lividus and Psammechinus miliaris are  
 19 submitted to the same environmental conditions in the Bay of Brest. The  
 20 relationship between seasonal changes in food source quality and their  
 21 gonad production was investigated in reproducing experimentally these  
 22 conditions. In a first stage two macroalgae (Palmaria palmata and Laminaria  
 23 digitata) were tested. P. miliaris showed a stronger preference for P. palmata  
 24 and over a year-long experiment both urchins progressively preferred P.  
 25 palmata. Seasonal variations in the chemical composition of P. palmaria

26 were observed in the Bay of Brest: total carbohydrates were important and  
27 the relative maximum (about 50%) was reached between February and  
28 August; the lipid level was low and had a relative maximum of about 1% in  
29 June and August. Total protein in P. palmaria was high compared to other  
30 seaweeds: the maximum value (25%) was observed in June, that was  
31 probably due to the maintenance of nitrogen nutrient in the bay.

32 In the second stage of the study, seasonal changes in biochemical  
33 components of ingestion and absorption of the two sea urchins were  
34 followed in the laboratory using a monospecific diet of P. palmaria. The  
35 patterns of total carbohydrates and lipid absorption were very similar for  
36 both sea urchin species. Carbohydrates were absorbed strongly and  
37 uniformly, year round. Lipid absorption mimicked the lipid nutrient pattern  
38 in the food source. Only changes in protein absorption varied slightly  
39 between the two urchin species. Protein absorption was maximal for both  
40 species in February and June, but the quantity of absorbed protein was  
41 significantly higher in P. miliaris than in P. lividus during February. This  
42 increase was concomitant with protein storage in the sea urchin gonads,  
43 which peaked in February for P. miliaris and in June for P. lividus. P.  
44 lividus had a higher gonad production efficiency, based on gonad yield. The  
45 comparison between in situ data and the experimental results suggests that  
46 an algal diet more nitrogenous than the in situ algal food source would  
47 benefit the herbivorous P. lividus, rather than the more omnivorous species  
48 P. miliaris. Although P. miliaris has been described as a species with large  
49 gonad production potential, P. lividus appears to be a more suitable species  
50 for echiniculture conditions.

51

52 Key words: sea urchin diet, *Palmaria palmata*. proximate composition,  
53 absorption efficiency, gonadal cycle.

54

## 55 1. Introduction

56

57 The sea urchins *Paracentrotus lividus* (Lamarck) and *Psammechinus miliaris*  
58 (Gmelin) are the two most common sea urchin species on the western coast  
59 of Brittany (France). Both species live in sheltered areas of intertidal and  
60 sublittoral zones. In an intertidal zone, *P. lividus* inhabits intertidal rock  
61 pools and *P. miliaris* lives under boulders; in subtidal zones, *P. lividus*  
62 occurs mainly on solid rocks or in seagrass meadows and has been observed  
63 on bottom sediments as diverse as gravels, heterogeneous sands or on maerl  
64 beds where it can cohabit with *P. miliaris* (Guillou et al., 2002). Both  
65 species have a commercial value. *P. lividus* populations have dramatically  
66 decreased on the northern coasts of Brittany because of destructive  
67 harvesting (Allain 1975, Southward and Southward, 1975). Although *P.*  
68 *miliaris* is smaller in size than *P. lividus*, it has a greater gonad production  
69 potential (Le Gall et al., 1989). Management of their populations could be  
70 improved by echiniculture.

71 Sea urchin biology, in general, has been well-studied all over the world,  
72 however studies of urchin populations in western Brittany are rare or  
73 incomplete for *P. lividus* (Allain, 1975, Dominique, 1973) and essentially  
74 for *P. miliaris* (Le Gall et al., 1989, 1990). Although both species have  
75 different areas of geographical distribution, they live in the Bay of Brest

76 under similar environmental conditions. Their different temperature optima  
77 can lead to different patterns of reproductive cycle in the present  
78 environment (Guillou, pers.obs.). Moreover, although they are inherently  
79 herbivorous, they can have different diet preferences (Boudouresque and  
80 Verlaque, 2001; Kelly and Cook, 2001). The purpose of this study is to use  
81 these specific differences to analyze the correlation between food quality  
82 and pattern of reproductive cycle in sea urchins.

83 In the first stage of our study, their dietary preferences among the  
84 macrophytes available in situ were tested by an experimental procedure. Sea  
85 urchins from the Bay of Brest were maintained in live under conditions as  
86 similar as possible to those of their natural habitat. A monospecific diet was  
87 desirable for the second stage of the study in which food ingestion rates and  
88 absorption rates were evaluated in terms of three major biochemical  
89 components: proteins, lipids and carbohydrates. These results were  
90 compared to the status of the sea urchins gonad production throughout a  
91 year-long experiment. Our approach combined simultaneous analyses of the  
92 seawater nutrients, the natural food source biochemistry and the absorption  
93 of different components by each species to explain changes in the gonad  
94 yield and composition during an annual cycle. The physiological responses  
95 of each species (food ingestion and absorption, reproductive growth) were  
96 also measured and compared with the goal of improving the culture of these  
97 two sea urchin populations.

98

99

## 100    **2. Materials and methods**

101

### 102    2.1 Sampling and maintenance

103

104    The reproductive cycle of adult Paracentrotus lividus and Psammechinus  
105    miliaris in the Bay of Brest was investigated from February 1997 to  
106    December 1998. The individuals were collected monthly by dredging or  
107    SCUBA divers from a site situated in the southern part of the Bay of Brest  
108    (Guillou et al., 2002) on substratum covered by maerl (a substrate composed  
109    of the living thalli of the calcareous red alga, Lithotamnion corallioides (P.  
110    and H. Crouan)). This substratum promotes the development of epiphytic  
111    macrophytes assemblages dominated by Rhodophyceae.

112    In the experimental study, P. lividus and P. miliaris individuals were  
113    collected by dredging in March 2000 in the same site. In the laboratory, the  
114    sea urchins were divided into three replicate groups consisting of 10  
115    individuals of each species, to measure feeding rates. Additionnal tanks  
116    maintained in the same experimental conditions were used for  
117    measurements of sea urchin gonad indices and biochemical analyses on the  
118    gonad tissues. A homogeneous size-class, representative of the dominant  
119    size-class of each population (Guillou et al., 2002), was selected: P. lividus:  
120    32-36 mm ( $34.3 \pm 1.8$ ) and P. miliaris 22-25 mm ( $24.1 \pm 1.5$ ). The sea urchin  
121    groups were placed in tanks (60 × 40 × 30cm) supplied with fresh running  
122    seawater from the Bay of Brest passed through on a sand-filter at  
123    temperatures which ranged from 9 °C in winter up to 17 °C in summer. A  
124    plastic grid of 2mm meshes on the evacuation exit of each tank prevented

125 the loss of algae or faeces. The photoperiod was adjusted weekly with a  
 126 timer by means of a set of neon tubes placed directly over the tanks (one 30-  
 127 watt tube per two tanks). Three replicate groups were used to measure  
 128 feeding rates.

129 A preliminary test for food preferences for the two species was  
 130 completed using: two green algae Cladophora rupestris (Linnaeus) Kützinger  
 131 and Enteromorpha ramulosa (Linnaeus), two red algae Palmaria palmata  
 132 (Linnaeus) O. Kuntze, Solieria chordalis (C. Agardh) J. and Plocamium  
 133 cartilagineum (Linnaeus) P. Dixon, and two brown algae Laminaria digitata  
 134 (Hudson) Lamouroux and Bifurcaria bifurcata (Ross). Three preferred algae  
 135 for the two sea urchins species were : P. palmata, S. chordalis and L.  
 136 digitata (Vachet and Guillou, pers. comm.). Because they were easier to  
 137 collect on a regular basis, P. palmata and L. digitata were used during the  
 138 long-term study. These algae were collected weekly from a site near the  
 139 laboratory facilities.

140

## 141 2.2 Feeding rates

142

### 143 2.2.1 First stage 2000-2001

144 In order to select which alga (Palmaria palmata or Laminaria digitata) was  
 145 preferred by the two urchins, algal ingestion rates of Paracentrotus lividus  
 146 and Psammechinus miliaris were recorded weekly in the laboratory from  
 147 March 2000 to July 2000 then from September 2000 to June 2001. Each  
 148 group of ten sea urchins was fed 10 g (WW, dried off in blotting paper) of  
 149 bits of P. palmata and 10 g of bits of L. digitata which were added

150 simultaneously in the tanks. Any food remaining after three days was  
 151 weighed and biomass was measured to the nearest 0.01 g (WW, dried off in  
 152 blotting paper). The ingested biomass (in g WW per urchin per day) was  
 153 calculated by subtraction. The loss of algal biomass during the time period  
 154 between feeding and collection has been estimated prior to the experiment  
 155 by weighing algae in three different tanks at different temperatures. The  
 156 algal loss was low,  $0.4 \pm 0.7$  % and  $1.4 \pm 1.3$  % at 12 and 17°C  
 157 respectively.. The 10 g algal ration added was always in excess of the  
 158 amount consumed both during and between the experiments. Tanks were  
 159 cleaned after each feeding session.

160

#### 161 2.2.2. Second stage 2001-2002

162 In the second part of the study, the ingestion rates and defaecation rates of  
 163 Paracentrotus lividus and Psammechinus miliaris, fed on the preferred alga  
 164 only, were recorded twice a month from October 2001 to August 2002.  
 165 Each group of ten sea urchins were fed with 15 g WW of the preferred alga.  
 166 All food offered, food remaining after 3 days and faeces collected through a  
 167 sieve were weighed. The faeces loss during the experiment was estimated  
 168 according to the procedure used for algae. This loss was  $2 \pm 3$  % and  $8.8 \pm$   
 169  $1.2$  % at 14 and 17°C respectively. For better precision, the biomasses were  
 170 expressed in dry weight to the nearest 1 mg. Because the offered biomass  
 171 was fresh and the water concentration varies seasonally in the alga, it was  
 172 converted to dry weight using the relationship between DW and WW  
 173 calculated at each feeding session. To do this, three samples of the alga were  
 174 first blotted dry in the paper, weighed, and then dried to constant weight



175 (48h at 60°C). The ratio of the wet weight /dry weight of these samples was  
176 calculated for the conversion. Algal biomass ingested and faeces produced  
177 and absorption, calculated as the difference between algal biomass ingested  
178 and faeces produced, were expressed in  $\text{mg DW. urchin}^{-1}.\text{day}^{-1}$ . Absorption  
179 rate was the ratio between absorption and the ingested biomass multiplied  
180 by 100.

181

### 182 2.3 Environmental parameters

183

184 Seawater samples were collected at a station close to the seawater intake  
185 that supplied the tanks in the laboratory and which was at less than 0.5  
186 nautical mile from the seaweed sampling site. Samples were collected  
187 weekly using the methods recommended by the French monitoring network  
188 in coastal environments (SOMLIT: <http://www.obs-vlfr.fr/somlit>).

189 Seawater was collected two meters below the surface at high tide and when  
190 the tide coefficient was  $70 \pm 10$ . Temperature was measured with a  
191 conductivity meter (LF 197). Seawater ammonium ( $\text{NH}_4^+$ ), nitrate ( $\text{NO}_3^-$ ),  
192 and nitrite ( $\text{NO}_2^-$ ) were measured according to the method described in  
193 Strickland and Parsons (1972), and modified for a Technicon autoanalyser  
194 with an accuracy of 5%.

195

196

### 197 2.4 Reproductive cycle

198

199 On each in situ sampling (from February 1997 to December 1998), 20  
200 individuals were brought back to the laboratory and dissected. Their gonads  
201 and tests were dried to constant weight (48h at 60°C). Gonad indices were  
202 calculated as the ratio of the dried gonad to the eviscerated test dry weight,  
203 and multiplied by 100.

204 Five times during the second stage of the experimental feeding experiment  
205 (24<sup>th</sup> October 2001, 21<sup>th</sup> December 2001, 5<sup>th</sup> February 2002, 7<sup>th</sup> June 2002,  
206 19<sup>th</sup> August 2002), five urchins of each species were isolated from the  
207 additional tanks to determine the gonad index according to the previous  
208 experimental protocol and to analyze the biochemical composition of the  
209 gonad.

210

## 211 2.5 Biochemical composition

212

213 The biochemical composition of the preferred alga, faeces and gonads were  
214 determined at the same time as gonad indices. The contents in  
215 carbohydrates, proteins and lipids of each compartment (alga, faeces and  
216 gonads) were determined. Three samples of algae and three samples of  
217 faeces from each urchin species were analyzed. Alga samples were rinsed  
218 and epiphytes removed before the analysis. Each sample of algae and faeces  
219 was divided into two parts. One part was weighed (wet weight) and then  
220 dried at 60°C to constant weight for estimation of the water content  
221 (difference between wet and dry weight). Ash content was determined on  
222 the dried tissue after combustion in a muffle furnace at 500°C for 4h. The  
223 second part of each sample was homogenized in distilled water using an

224 Ultra turax and this homogenate was used for the biochemical analyses.  
225 Carbohydrates were analysed using the Dubois procedure (Dubois *et al.*,  
226 1956). Nitrogen was determined by the total Kjeldahl method (TKN)  
227 (protein content =  $6.25 \times \text{TKN}$ ) (Indergaard and Minsaas, 1991). Total lipid  
228 content was determined gravimetrically using the Bligh and Dyer method  
229 (1959).

230 For the gonad analyses four sea urchins were dissected and their gonads  
231 collected and homogenized with the Ultra turax. This homogenate was  
232 divided in four parts: the first split was used for water content  
233 determinations (drying at 60°C to constant weight). The dried material was  
234 then combusted at 500°C for 4h to determine the ash content of gonads. The  
235 remaining 3 splits were used for measuring the levels of carbohydrates,  
236 proteins and lipids using the techniques of Dubois et al. (1956), Lowry et  
237 al.(1951) and Bligh and Dyer (1959), respectively.

238 The proximate organic composition of each compartment was determined  
239 using the ash-free dry weight (AFDW). From these data, ingestion rates in  
240 terms of organic components, (carbohydrates, proteins and lipids) were  
241 expressed in  $\text{mg DW.urchin}^{-1}.\text{day}^{-1}$  for each nutrient.

242 The quantity of the ingested component was equal to the percentage of this  
243 component present in the alga sample at any given period multiplied by the  
244 quantity of alga ingested by the sea urchin over the same time. The quantity  
245 of excreted component was a function of the percentage of this material in  
246 the faeces and the quantity of faeces produced by the sea urchin. The  
247 quantity of component absorbed by the organism was the difference  
248 between the quantity ingested and the quantities excreted.

249 The chemical composition of the gonads was corrected by the gonad index  
250 at the time of sampling in order to take into account the changes in gonad  
251 weight over the length of the experiment. The index was calculated from the  
252 percentage of the organic component in the gonad at a given time multiplied  
253 by the gonad index at the same sampling.

254

## 255 2.6 Statistics

256 Changes in ingestion and defaecation rates, gonad index, quantities of  
257 ingested components (carbohydrates, proteins, lipids), of absorbed  
258 components and chemical composition of the gonad, were tested for each  
259 sea urchin species with a one-way analysis of variance (ANOVA) ( $P <$   
260  $0.05$ ) with the least significant difference test once the homogeneity of  
261 variance had been tested. The gonad index of experimental and control  
262 animals were arcsine-transformed.

263 All analyses were done with the statistical software STATGRAPHICS 4.

264

265

## 266 **3 Results**

267

### 268 3.1 Environmental variations

269

270 Ammonium ( $\text{NH}_4^+$ ), nitrate ( $\text{NO}_3^-$ ), and nitrite ( $\text{NO}_2^-$ ) levels increased  
271 beginning in October 2001 (Fig.1a). The main peak of ammonium was  
272 observed at the end of October ( $2.4 \mu\text{M}$ ) followed by a nitrite peak at the  
273 end of November ( $0.75 \mu\text{M}$ ) and a nitrate concentration peak in mid-

274 February 2002 (23.6  $\mu\text{M}$ ). Then nitrates decreased in March when  
 275 chlorophyll *a* showed a small peak (Fig. 1a and b). Nitrites and nitrates  
 276 dropped to very low levels in March and May respectively, and stayed low  
 277 until September during the temperature maximum (Fig. 1a).

278 Ammonium reached its lowest levels from March to the end of June  
 279 followed by a new peak at the beginning of August. Successive peaks of  
 280 chlorophyll *a* occurred from mid-May to the end of August (Fig. 1b).

281

### 282 3.2 Reproductive cycle

283

284 Field data obtained in 1997 and 1998 on the Paracentrotus lividus  
 285 reproductive cycle in the Bay of Brest indicated that the time when  
 286 spawning started, marked by a drop in the GI, differed between years (Fig.  
 287 2a). In 1997, the GI reached a maximum in May (GI=7) and then decreased  
 288 sharply, indicating a short spawning period. In contrast during 1998, the GI  
 289 decrease was small during winter and spring. Each year, the minimum GI  
 290 values were observed in June and followed by a rapid increase. Spawning of  
 291 Psammechinus miliaris occurred from early March to mid-June in 1997 and  
 292 from mid-April to mid-June in 1998 (Fig. 2b). The GI reached maximum  
 293 values of 12 and 8 respectively, and a minimum value of 2. This low level  
 294 reflecting the resting stage remained steady during about 3 months.

295 For both species P. lividus and P. miliaris, the changes in gonad indices (GI)  
 296 under experimental conditions confirmed the seasonal variations (Fig. 2a  
 297 and b). For P. lividus, the GI increased from October 2001 (GI=2) to June  
 298 2002 (GI=8). In August, the GI value was still high. Comparison with the

field data suggested the spawning event in the experimental study would be around the maximum GI value (8) observed in June. After the onset of spawning, which cannot be precisely defined here, the GI might drop to a low level located in mid-June in both sets of field data (1997 and 1998). Spawning marks were observed visually in the laboratory tanks during this time. Thus, the GI estimated in August would be during the recovery stage of the gonad, as the post-spawning stage, or resting stage, was very short in the field confirming the previous studies on P. lividus (Byrne, 1990; Spirlet et al., 1998). For P. miliaris, GI values increased significantly from December 2001 (GI=5) and reached the highest value (15) in February 2002, after which the GI value decreased and reached a minimum level in August 2002 (GI=2.6). The high level observed in February compared to the field data situated the onset of spawning event close to February. As for P. lividus the two sets of 1997 and 1998 field data indicated that the end of spawning took place in June. The very low value measured in August 2002, and similar to the field observations, suggests the gonads were in resting stage. The field and experimental observations indicate that spawning occurred earlier in P. miliaris than in P. lividus.

317

### 318 3.3 Feeding preference

319

The feeding rates on Palmaria palmata and Laminaria digitata for the two urchin species Paracentrotus lividus and Psammechinus miliaris from March 2000 to June 2001 are presented in the figure 3 using units of g WW. urchin<sup>-1</sup>.day<sup>-1</sup>. With respect to P. lividus (Fig. 3a) three feeding rate trends were

324 observed: from March 2000 to July 2000 (except for April 2000), sea  
 325 urchins ingested quantities significantly larger of L. digitata than P. palmata,  
 326 then from September 2000 to June 2001, the ingestion of L. digitata and P.  
 327 palmata did not differ significantly, and finally, from May 2001 to June  
 328 2001, the ingested biomass of P. palmata were higher than those of L.  
 329 digitata ones ( $P < 0.05$ ).

330 For P. miliaris, (Fig.3b), two stages could be distinguished: from March  
 331 2000 to September 2000, the feeding rates on L. digitata and P. palmata  
 332 were not significantly different, and in the second stage, from October to  
 333 June 2001, more P. palmata was ingested than L. digitata ( $P < 0.05$ ). The  
 334 ingestion rate of P. palmata increased significantly during this last period.  
 335 This increase coincided with a decreasing consumption of L. digitata over  
 336 the same period.

337 Finally the both species presented a similar pattern with a higher attraction  
 338 for P. palmata with time .

339

#### 340 3.4 Ingested and defaecated biomasses during 2001-2002

341

342 Based on the previous results, Palmaria palmata was used for the second  
 343 part of the study. Samples were collected in Dellec Cove, near the seawater  
 344 sampling station. In both species, changes in ingestion and defaecation rates  
 345 had a similar pattern, with more pronounced variations in Psammechinus  
 346 miliaris than in Paracentrotus lividus (Fig. 4a and b). A decrease in  
 347 ingestion rate was observed from February ( $60 \text{ mg DW urchin}^{-1} \text{ d}^{-1}$ ) to April  
 348 ( $30 \text{ mg DW urchin}^{-1} \text{ d}^{-1}$ ) in P. lividus, and from January ( $79 \text{ mg DW urchin}^{-1}$

349  $^1 \text{ d}^{-1}$ ) to April (20.5 mg DW urchin $^{-1} \text{ d}^{-1}$ ) in P. miliaris. After April, ingestion  
 350 rates increased through June (50 and 68 mg DW urchin $^{-1} \text{ d}^{-1}$  for P. lividus  
 351 and P. miliaris respectively) and remained high through summer.

352 The pattern of defaecation followed that of ingestion with minima observed  
 353 in April. Changes were significantly more pronounced for P. miliaris than  
 354 for P. lividus.

355

### 356 3.5 Biochemical composition of Palmaria palmata

357

358 Biochemical analyses done on Palmaria palmata five times during the year  
 359 showed seasonal changes in organic component levels (Fig. 5).

360 Carbohydrates increased significantly from October (40.4 % AFDW) to  
 361 December (53.2 %) then remained constant until August ( $P < 0.05$ ).

362 Proteins increased significantly from December (12.4 %) to February (24.4  
 363 %) and then decreased from June to August (13.7 %). The maximum level  
 364 of proteins in P. palmata was measured in February and June.

365 Lipids increased significantly from February (0.4 %) to June (1.1 %) and  
 366 reached their maximum value in August (1.3 %).

367

### 368 3.6 Quantity of ingested nutrients

369

370 The estimated ingestion of carbohydrates remained constant for  
 371 Paracentrotus lividus throughout the annual cycle, about 20 mg DW. urchin $^{-1}$   
 372  $^1 \text{ day}^{-1}$  ( $P > 0.05$ ) (Fig. 6a). For Psammechinus miliaris, the quantity of  
 373 ingested carbohydrates increased significantly from October (19.6 mg DW.



374 urchin<sup>-1</sup>.day<sup>-1</sup>) to December (26.8 mg DW. urchin<sup>-1</sup>.day<sup>-1</sup>) and reached its  
 375 maximum level in February and June (29.5 mg DW. urchin<sup>-1</sup>.day<sup>-1</sup>) (Fig.  
 376 6b). Then it decreased from June to August (26.1 mg DW. urchin<sup>-1</sup>.day<sup>-1</sup>).

377 The estimated quantity of proteins ingested by P. lividus and P. miliaris,  
 378 increased significantly from October (6.3 mg and 6 mg DW. urchin<sup>-1</sup>.day<sup>-1</sup>,  
 379 respectively) to February (9.6 mg and 14.3 mg DW. urchin<sup>-1</sup>.day<sup>-1</sup>,  
 380 respectively). However, in P. lividus the maximum level occurred in June  
 381 (12.1 mg DW. urchin<sup>-1</sup>.day<sup>-1</sup>) ( $P < 0.05$ ), while in P. miliaris it was  
 382 observed in both February (14.3 mg DW. urchin<sup>-1</sup>.day<sup>-1</sup>) and June samples  
 383 (15.7 mg DW. urchin<sup>-1</sup>.day<sup>-1</sup>) which were not significantly different. In both  
 384 species the quantity of proteins ingested decreased significantly between  
 385 June and August.

386 With respect to the lipids, the estimated quantity ingested by each species  
 387 increased significantly between February (0.17 and 0.18 mg DW. urchin<sup>-1</sup>.  
 388 day<sup>-1</sup>, respectively) and June (0.53 and 0.68 mg DW. urchin<sup>-1</sup>.day<sup>-1</sup>,  
 389 respectively). Maximum levels of lipids were ingested in June and August.

390

### 391 3.7 Total absorption rate and quantity of absorbed components

392

393 The total absorption rate was high for both species (Fig. 4). In  
 394 Psammechinus miliaris a period of low absorption occurred in May ( $60.1 \pm$   
 395  $6.36\%$ ) between two periods of high, but significantly different, absorption  
 396 rates, the first from October to the end of April ( $82.1 \pm 5.5\%$ ) and the  
 397 second from the mid-June to the end of August ( $77.6 \pm 4.9\%$ ). In  
 398 Paracentrotus lividus the absorption rate was homogeneous over the year

399 (87.6  $\pm$  3 %) and was significantly higher than even the high absorption rate  
 400 periods of P. miliaris ( $P < 0.05$ ). With respect to the different components,  
 401 the absorption of carbohydrates was significantly higher in P. miliaris than  
 402 in P. lividus (97  $\pm$  1% versus 86  $\pm$  7% in). The protein absorption did not  
 403 vary significantly between the two species (78  $\pm$  9% and 80.5  $\pm$  7% for P.  
 404 miliaris and P. lividus respectively).

405 The amount of an absorbed biochemical component was considered relative  
 406 to the ingested and defaecated biomass of the same component (Fig. 7). The  
 407 quantity of absorbed carbohydrates was not significantly different during the  
 408 annual cycle for each species but was significantly different ( $P < 0.05$ )  
 409 between both species with 20.7  $\pm$  1 and 26.4  $\pm$  4 mg DW.urchin<sup>-1</sup>.day<sup>-1</sup> for P.  
 410 lividus and P. miliaris respectively. Both species exhibited similar changes  
 411 in the absorption of proteins. The quantity of absorbed proteins increased  
 412 significantly from October (4.9 and 4.8 mg DW.urchin<sup>-1</sup>.day<sup>-1</sup> for P. lividus  
 413 and P. miliaris respectively) to February (7.12 and 9.9 mg DW.urchin<sup>-1</sup>.day<sup>-1</sup>  
 414 1) and then from February to June (11.2 and 13.8 mg DW.urchin<sup>-1</sup>.day<sup>-1</sup>).  
 415 This increase was followed by a decrease from June to August (4.3 and 5.7  
 416 mg) ( $P < 0.05$ ).

417 For both P. lividus and P. miliaris, the absorption of lipids was only  
 418 quantifiable in June and August (0.37 and 0.46 mg DW.urchin<sup>-1</sup>.day<sup>-1</sup> in P.  
 419 lividus and P. miliaris, respectively). because of the scarcity of this  
 420 component in the alga.

421

422 3.8 Biochemical composition of the gonad

423

424 The quantity of carbohydrates in the gonad increased significantly for both  
 425 species, from December to February and then decreased from February to  
 426 June ( $P < 0.05$ ) (Fig. 8).

427 The protein content in Paracentrotus lividus gonads increased steadily and  
 428 significantly from October and reached its maximum level in June ( $P <$   
 429  $0.05$ ); it decreased between June and August, but remained superior to  
 430 October and December values. For Psammechinus miliaris, the quantity of  
 431 proteins in the gonads increased significantly from December to February  
 432 then decreased steadily and significantly to August. The level in August was  
 433 lower than that in October ( $P < 0.05$ ).

434 The quantity of lipids in gonad samples increased significantly for P. lividus  
 435 from December to February, decreasing thereafter into August. For P.  
 436 miliaris, an important significant increase was observed between December  
 437 and February, followed by successive significant decreases in both June and  
 438 August ( $P < 0.05$ ).

439

440

## 441 **4. Discussion**

442

### 443 **4.1 Palmaria palmata as a nutritional source**

444

445 One of our first objectives was to determine the preferred alga by the  
 446 two sea urchin species in order to use a monospecific, natural diet for  
 447 subsequent experiments. A previous study (Vachet, unpublished) suggested  
 448 the sea urchins had a preference for two algae already used commonly in

449 echiniculture: Palmaria palmata and Laminaria sp. (Basuyaux and Blin,  
450 1998; Kelly, 2001, Spirlet et al., 2000). In the present study, sea urchins  
451 were fed P. palmata and L. digitata for more than one year. Analysis of the  
452 results showed that, in the short term (6 months), there was a variable  
453 consumption rate of the two algae, by the sea urchin species. Over longer  
454 time periods, there was a progressively greater consumption of P. palmata  
455 by both urchin species. In this first experiment, this change in feeding  
456 preference was not directly correlated to changes in alga composition or in  
457 sea-urchin maturity as the feeding response during the period of intense  
458 modifications in algae and in sea-urchin gonads (April-June) was  
459 significantly different between 2000 and 2001. Lemire and Himmelman  
460 (1996) have classified different algae according to their ability to support  
461 somatic and gonadic growth (using hierarchical cluster analysis), and reported  
462 that both these algae contributed strongly to the fitness of another urchin  
463 species, Strongylocentrotus droebachiensis. Vadas et al. (2000) in a similar  
464 study, also concluded that P. palmata among four species of preferred  
465 macroalgae “induced the quickest and highest” enhancement in gonad index  
466 values. The improvement in gonad yield has been credited to the high  
467 protein levels measured in this alga (Fleurence, 1999 and Martinez and  
468 Rico, 2002), an explanation discussed by other investigators (see review  
469 Morgan et al., 1980 and Hagen Rødde et al. 2004). L. digitata contains a low  
470 proportion of protein and a relatively high proportion of complex  
471 carbohydrates (Otero-Villanueva et al., 2004) that can explain the poorer sea  
472 urchin ingestion, absorption and assimilation efficiencies.

473 Our study showed an increase in total protein in the alga, P. palmata  
 474 between October and December, and maximum values were reached in  
 475 February and June (24.4 % AFDW). These values were close to the  
 476 maximal values reported from other studies in Brittany (about 25% from  
 477 March to May in the southern part (Galland-Irmouli et al., 1999) and 22 to  
 478 20.4% between February and April in the northern part (Rouxel et al.,  
 479 2001)) and were superior to values reported from the northern Spanish alga  
 480 populations (18 % between March and May (Martinez and Rico, 2002)).  
 481 The main difference between all these populations was the maintenance of a  
 482 high protein level during June in P. palmata from the Bay of Brest, while the  
 483 protein level decreased to 10 % in other populations along the coast of  
 484 Brittany and declined to 2 % at the Spanish sites. The maintenance of a high  
 485 protein content in P. palmata was probably related to the seawater nitrate  
 486 concentration. Nitrate is the most available N source and is the main  
 487 inorganic nutrient involved in algal nutrition (Chapman and Craige, 1977).  
 488 A rapid increase in protein contents of P. palmata follows high  
 489 concentrations of seawater nitrate (Morgan and Simpson, 1981). In our  
 490 study, the increase was concomitant with the increase in seawater  $\text{NO}_3^-$  and  
 491  $\text{NO}_2^-$  concentrations and the maximum protein content occurred during the  
 492 peak of  $\text{NO}_3^-$ . The overall seawater nitrogen concentrations in the Bay of  
 493 Brest (maximum  $\text{NO}_3^- + \text{NO}_2^-$ : 24  $\mu\text{M}$ ) was higher compared to those on the  
 494 Spanish coast (9  $\mu\text{M}$ ).

495 P. palmata in our study remained very rich in proteins even in June.  
 496 These proteins serve as a reserve source used for growth, maintenance and  
 497 reproduction by the alga. In Brittany, the reproductive stage of P. palmata

occurs during winter and the maximum growth rate, during winter and spring (Le Gall, 2002). Thus in June the protein content should have been low in the alga as it is the case in the Spanish coast, except if a nitrogen source was still present in the seawater., Two indices suggests the higher level of nitrogen in the Bay of Brest; the first is the presence of low but not insignificant concentrations of  $\text{NH}_4^+$  which can also be utilized by the algae to contribute to the maintenance of growth (Martinez and Rico, 2002). The second is the occurrence of successive peaks of chlorophyll *a*, corresponding to phytoplankton blooms, from May to the end of August. These summer peaks of low intensity typical of the Bay of Brest ecosystem (<http://www.obs-vlfr.fr/somlit>) suggest sufficient nutrients were present to support bloom conditions, which could benefit the macroalgae also.

510

#### 4.2 Changes in ingestion and defaecation rates

512

In the two sea urchin species, monthly variations were observed for both ingestion and defaecation rates. The possible loss in alga and faeces biomass during the experiment was too low to explain the main changes. The difference in timing for the start of an ingestion rate decrease (in January for *P. miliaris* and in February for *P. lividus*) may also be related to the relative stage of maturity in each species. During 2002, the highest GI reported here, and corroborated by the earlier field data (Fig. 2), showed that the maturity stage occurred earlier in *P. miliaris* than in *P. lividus* with the bay of Brest environmental conditions. Some previous studies have shown that echinoid feeding rates decrease before spawning (Fuji, 1967, De Ridder

523 and Lawrence, 1982). The reason for this phenomenon may be  
 524 physiological or due to the gonad size increase into the coelomic space  
 525 during the gametogenesis. The first hypothesis is plausible for both species,  
 526 but the second only concerns P. miliaris, since the P. lividus GI was high in  
 527 April when feeding activity increased again. In both species, the increase in  
 528 food consumption was concomitant with a water temperature increase in  
 529 mid-April, suggesting temperature can control the sea urchin feeding rates  
 530 also (see review Lares and Mc Clintock, 1991).

531 The defaecation rate changes in both species mimicked, in general, changes  
 532 in ingestion rates. The total nutrient absorption rates were high (mean  
 533 annual values of 78% and 62 % for P. lividus and P. miliaris respectively)  
 534 but not superior to the values observed in P. lividus by Frantzis and  
 535 Grémare (1992), often above 80%. P. miliaris presented absorption rates  
 536 significantly lower and seasonal changes in ingestion and defaecation rates  
 537 more pronounced than P. lividus For P. miliaris, total nutrient absorption  
 538 was significantly lower after the spawning event, than between October and  
 539 April during the gametogenesis stage. This process is probably related to  
 540 progressive increase of reserve storage for gametogenesis.

541

542 4.3 Changes in nutrient absorption rate: connection with the proximate  
 543 composition of food and gonad

544

545 Absolute changes in absorption rate differed for each nutrient, but the  
 546 patterns were very similar for both species. The carbohydrates were  
 547 absorbed uniformly throughout the year, in contrast to the absorption of

548 proteins and lipids, which changed seasonally. The absorption of proteins  
549 significantly increased from October to June, and then decreased from June  
550 to August when the absorption of lipids increased. These changes in sea  
551 urchin nutrient absorption were linked to several factors: the total  
552 concentration of the nutrient in the food, the specific composition of lipids,  
553 carbohydrates and proteins, the physiological requirements of the sea urchin  
554 for a particular nutrient, and the digestive characteristics of the sea urchin,  
555 (especially its enzymatic equipment). Without data on changes in specific  
556 composition of the nutrients and their digestibility in the sea urchins, this  
557 discussion was only based on the relationship between the proximate  
558 organic composition of the alga and its absorption by the sea urchins with a  
559 particular attention to the gonad production.

560 For the two sea urchin species in our study, carbohydrate absorption was not  
561 affected by diet as has been previously described in Watts et al., (1998) for  
562 Lytechinus variegatus (L). In our study, the carbohydrate absorption did not  
563 vary during the year-long experiment, even though this component  
564 increased significantly in Palmaria palmata from October to December. The  
565 carbohydrate absorption rate strongly suggest that these sea urchins were  
566 efficient in digesting the available carbohydrates. However, overall lower  
567 carbohydrate absorption recorded for *P. miliaris* suggest that *P. lividus* has  
568 better enzymatic conditions for digesting the insoluble carbohydrate fraction  
569 (which can represent about 55% of the dry weight of P. palmata) (Lahaye,  
570 1991; Hagen Rødde et al., 2004). Total carbohydrate absorption was  
571 probably not affected by physiological demand for reproduction because the  
572 maximum need in this component (essentially as glycogen, Monteiro-



573 Torreiro and Garcia-Martinez, 2003) would have been between February  
574 and June for P. lividus, and December and February June for P. miliaris.  
575 Lipid absorption was only observed in June and August when their levels  
576 were maximal in P. palmata. With a total lipid content of more than 1% in  
577 summer, P. palmata in the Bay of Brest have a relatively high lipid  
578 concentration (Sanchez-Machado et al., 2004). There was no significant  
579 difference in the mean quantity of lipids absorbed by the two sea urchins  
580 during this period. Their absorption reflects the significant increase of lipids  
581 in the food source and could not be linked to reproductive needs: the  
582 maximum gonad demand for this nutrient was in February for both sea  
583 urchin species.

584 The protein level in P. palmata increased from October to February. This  
585 increase was followed by the increase of ingested proteins from October to  
586 February for P. miliaris and from October to June for P. lividus. In both  
587 species, the maximum level of absorbed protein was observed in June. From  
588 February, the quantity of absorbed protein was significantly higher in P.  
589 miliaris than in P. lividus. These observations attest to a physiological  
590 relationship between the increase in the protein absorption and reproduction,  
591 the gonad growth phase being earlier in P. miliaris than in P. lividus. Protein  
592 is the major component of P. lividus and P. miliaris gonads (Monteiro-  
593 Torreiro and Garcia-Martinez, 2003) and the need for this nutrient increases  
594 strongly before spawning (Fenaux et al., 1977; Fernandez, 1998, Monteiro-  
595 Torreiro and Garcia-Martinez, 2003). In our study, this requirement was  
596 highest in February for P. miliaris and in June for P. lividus and would have  
597 been supported by the high protein content in P. palmata production during

598 the same period. The protein conversion from ingested food to gonad  
 599 biomass is known to be rapid (Fernandez, 1996) and suggests that gonadal  
 600 growth cannot be effective when only protein reserves are available. The  
 601 organism needs the protein-rich food also.

602 The relationship between gonad yield and protein content in algae (Lowe  
 603 and Lawrence, 1976; Larson et al., 1980; Vadas et al., 2000) or in prepared  
 604 feeds (see review Pearce et al., 2003) is well-documented. Comparing the  
 605 GI obtained experimentally with the monospecific P. palmata diet and the  
 606 GI observed in the field suggested that this protein-rich alga enhances the  
 607 gonad yield in P. lividus. The results were less clear in P. miliaris. This  
 608 species is known to be more omnivorous than P. lividus (*op.cit.*) and under  
 609 natural conditions, P. miliaris feeds on algae and large numbers of  
 610 encrusting intertidal organisms such as mussels or barnacles (Kelly and  
 611 Cook, 2001), increasing its protein input.

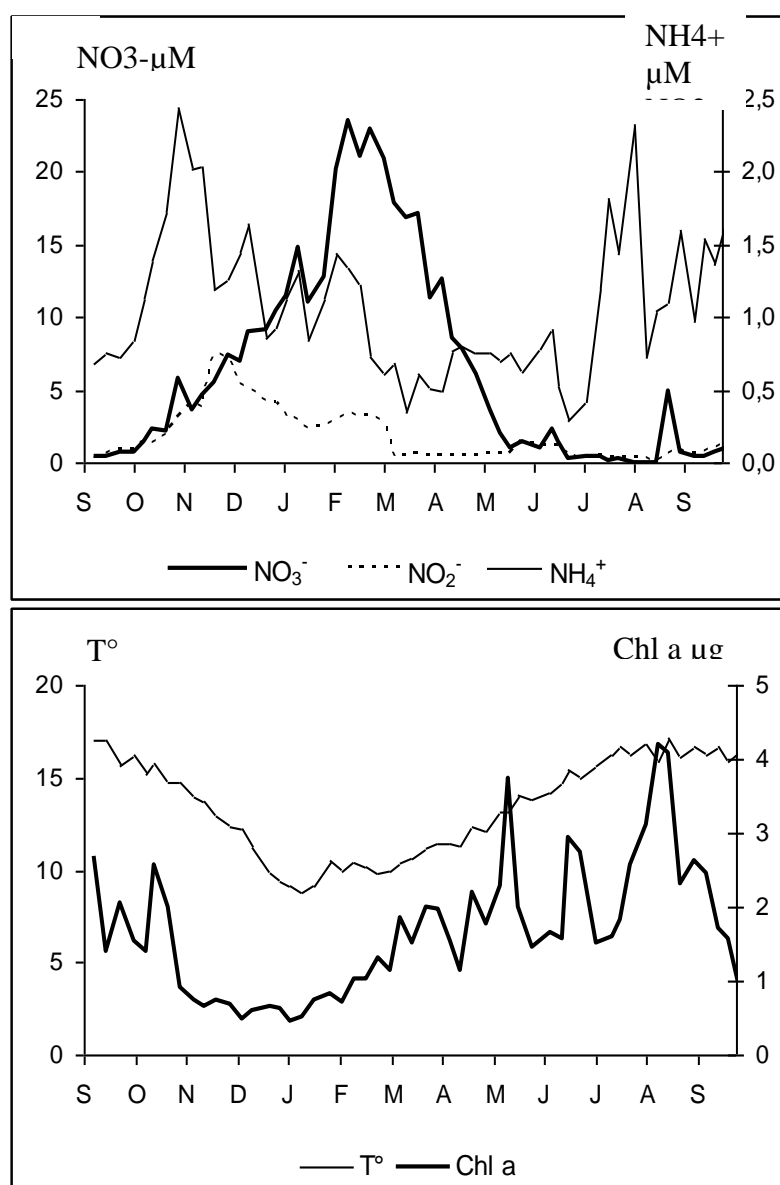
612 Our experimental results showing the stronger preference of P. miliaris for  
 613 the more protein-rich alga P. palmata (as compared to P. lividus) is  
 614 consistent with the possibility that P. miliaris has a higher protein  
 615 requirement. Higher protein ingestion may also explain the higher in situ P.  
 616 miliaris GI values as compared to those of P. lividus (Le Gall, 1989; Kelly,  
 617 2000; this study). The enhanced gonad index in P. lividus when fed a  
 618 monospecific high protein diet suggests that the optimum protein level  
 619 (Akiyama et al., 2001) to maximize P. lividus gonad production is not  
 620 reached under natural conditions, compared to P. miliaris. A protein-rich  
 621 algal diet, atypical for P. lividus, could favour gonad growth in this species,  
 622 whereas P. miliaris can utilise food of animal origin under natural

623 conditions. The quantity of ingested and absorbed nutriments per urchin per  
624 day related to the sea urchin test biomass was higher in P. miliaris than in P.  
625 lividus. However, the maximum gonad biomass recorded in 2002 from P.  
626 miliaris (0.45g DWW) remained low compared to the maximum gonad  
627 biomass from P. lividus (0.70g DWW). In the same way, the conversion  
628 efficiency of food to gonadal production at a mature stage (ratio of ingestion  
629 rate to gonad growth rate) is better for *P. lividus* than for P. miliaris (20%  
630 and 9% respectively). Under echiniculture conditions, gonad production  
631 enhancement by protein input from natural food sources is likely to be more  
632 productive for P. lividus than P. miliaris.

633

634

## 635 LEGENDS

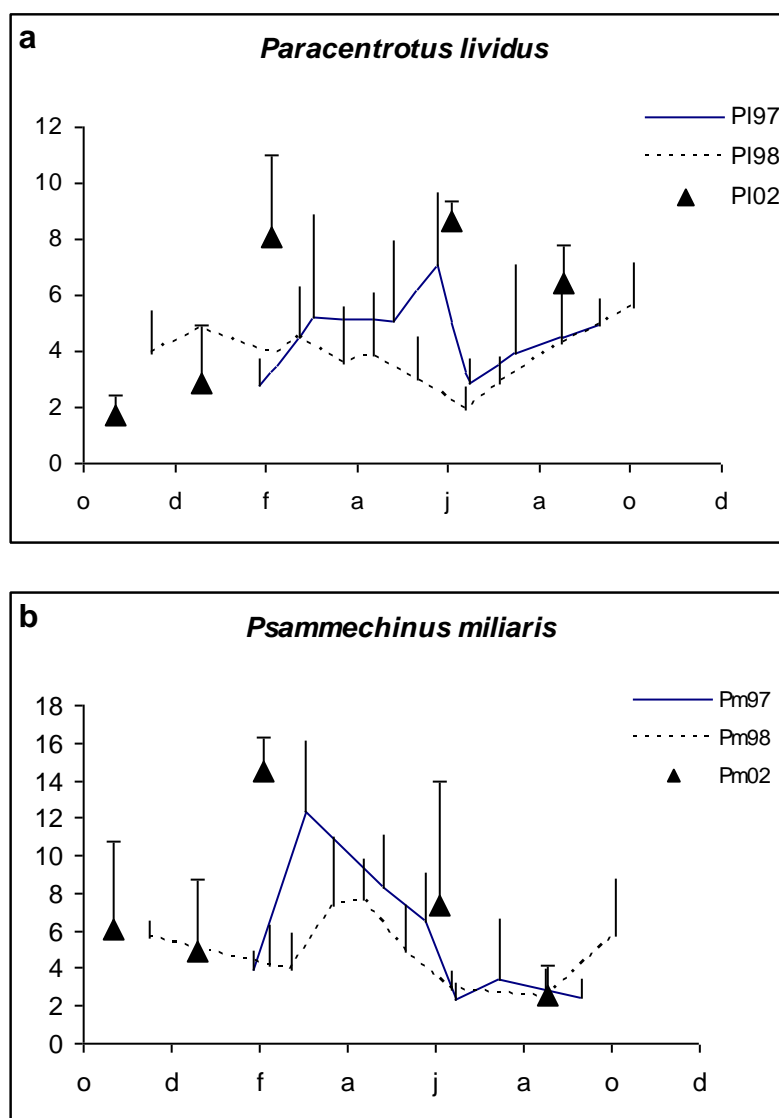


636

637 Fig.1. Seasonal changes in the seawater parameters in the Bay of Brest. from

638 September 2001 to October 2002 : a : ammonium, nitrite and nitrate; b :

639 temperature and chlorophyll *a*



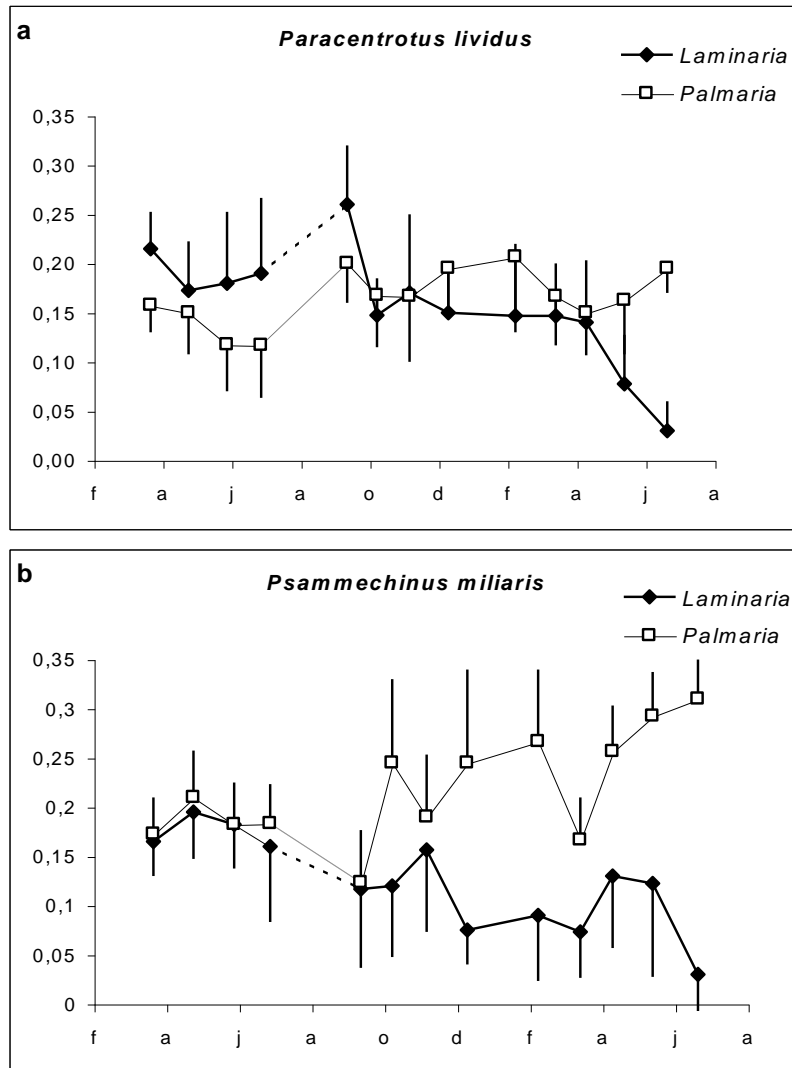
640

641 Fig. 2 Gonad indices (in % of dry weight) during the experiment (black

642 triangle +SD) compared to the IG seasonal changes recorded in 1997 and

643 1998 from *in situ* populations

644



645

646 Fig. 3 Seasonal changes in the biomass of Laminaria digitata and Palmaria647 palmata ingested by the sea urchins (in g WW d<sup>-1</sup> urchin<sup>-1</sup>) ( $\pm$ SD) from648 March 2000 to March 2001; a : Paracentrotus lividus; b : Psammechinus649 miliaris.

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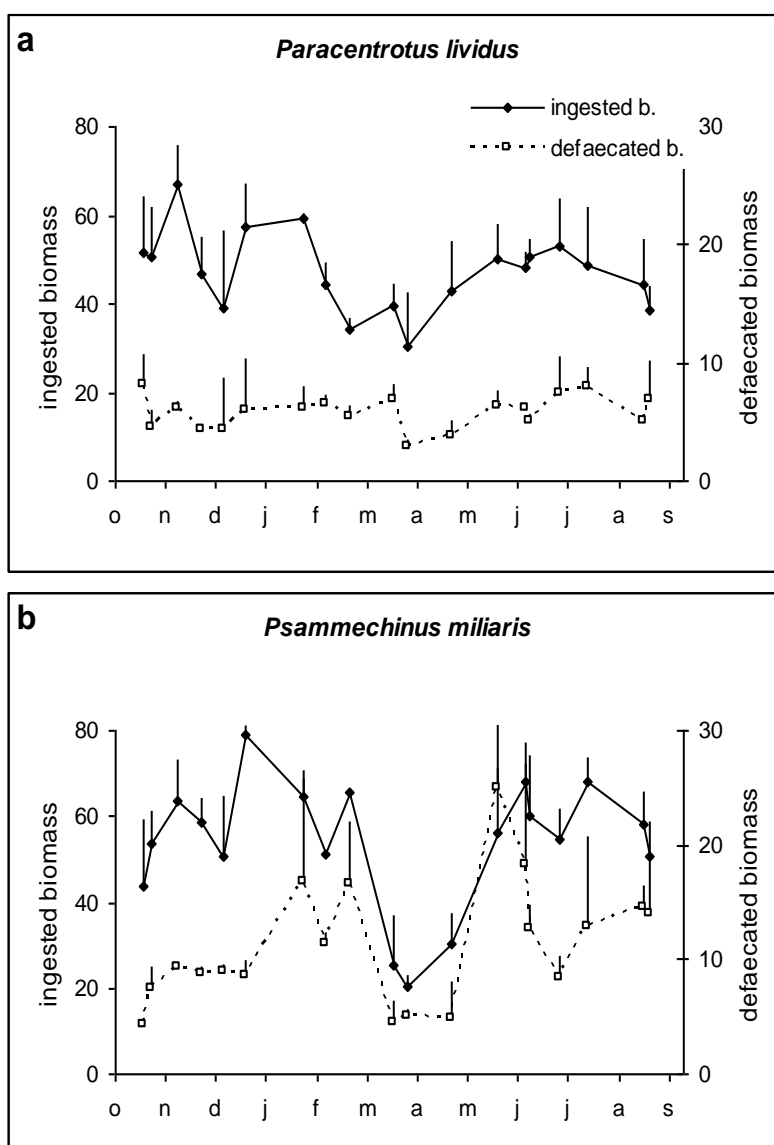
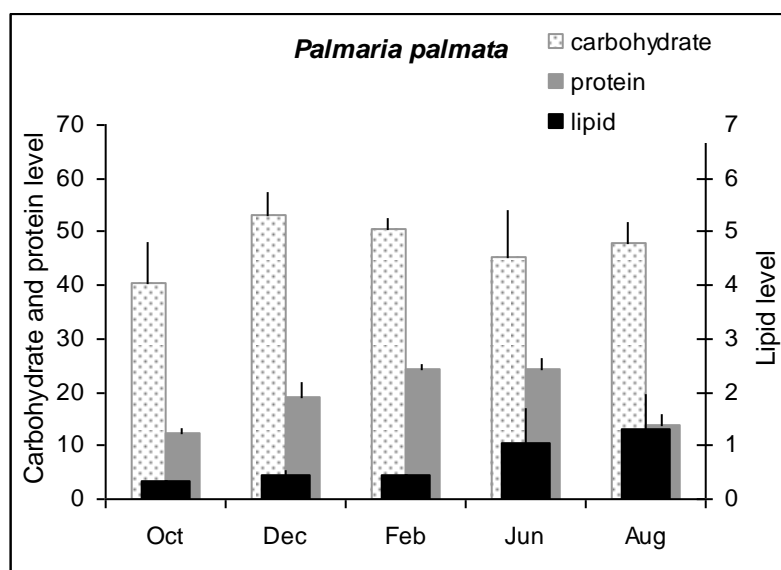


Fig. 4 Seasonal changes in the ingestion and defaecation rate of the sea urchins fed *Palmaria palmata* (in mg DW d<sup>-1</sup> urchin<sup>-1</sup>) (+SD) from October 2001 to August 2002. a : *Paracentrotus lividus*; b : *Psammechinus miliaris*.



671

672 Fig. 5 Seasonal changes in the proximate organic composition of Palmaria673 palmata (in % of DW) (+SD)

674



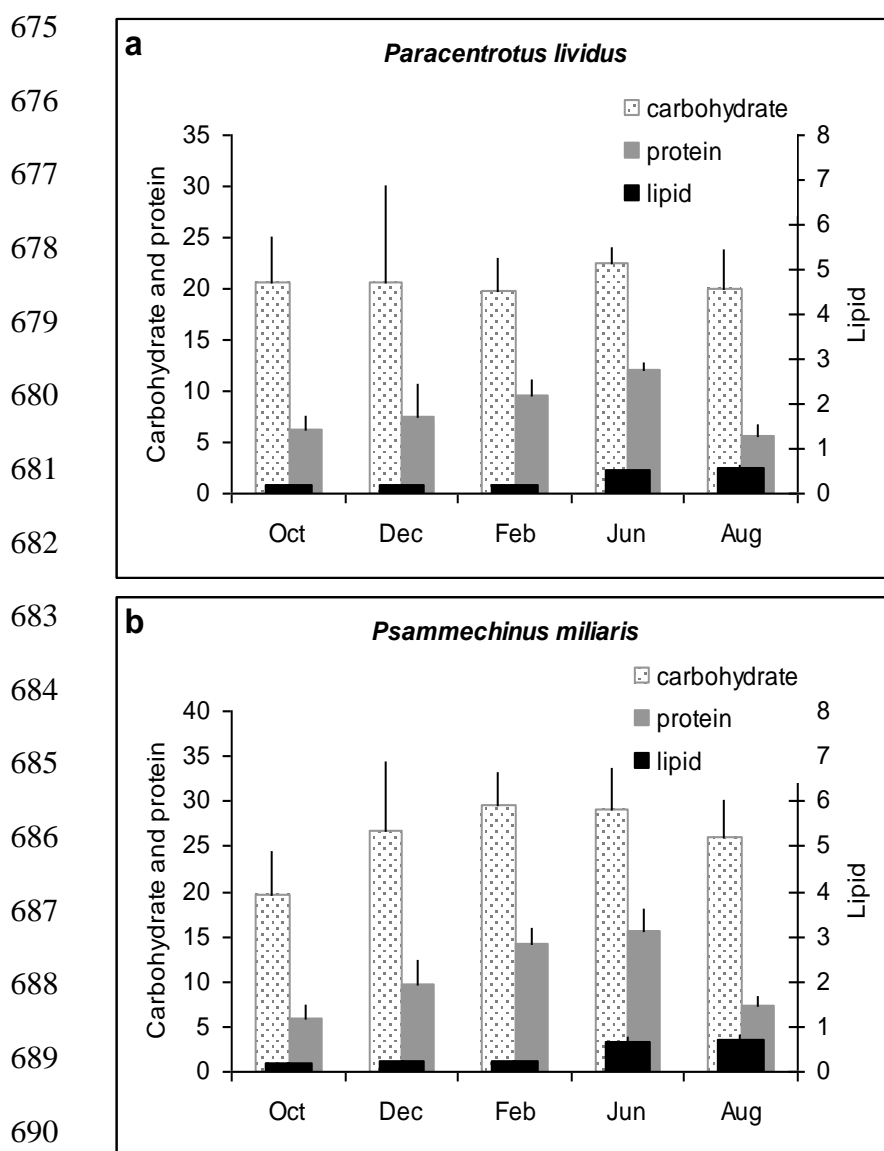


Fig. 6 Seasonal changes in the ingestion rate of the sea urchins fed *Palmaria palmata* in term of proteins, carbohydrates and lipids (in mg DW d<sup>-1</sup> urchin<sup>-1</sup>) (+SD). a : *Paracentrotus lividus*; b : *Psammechinus miliaris*.

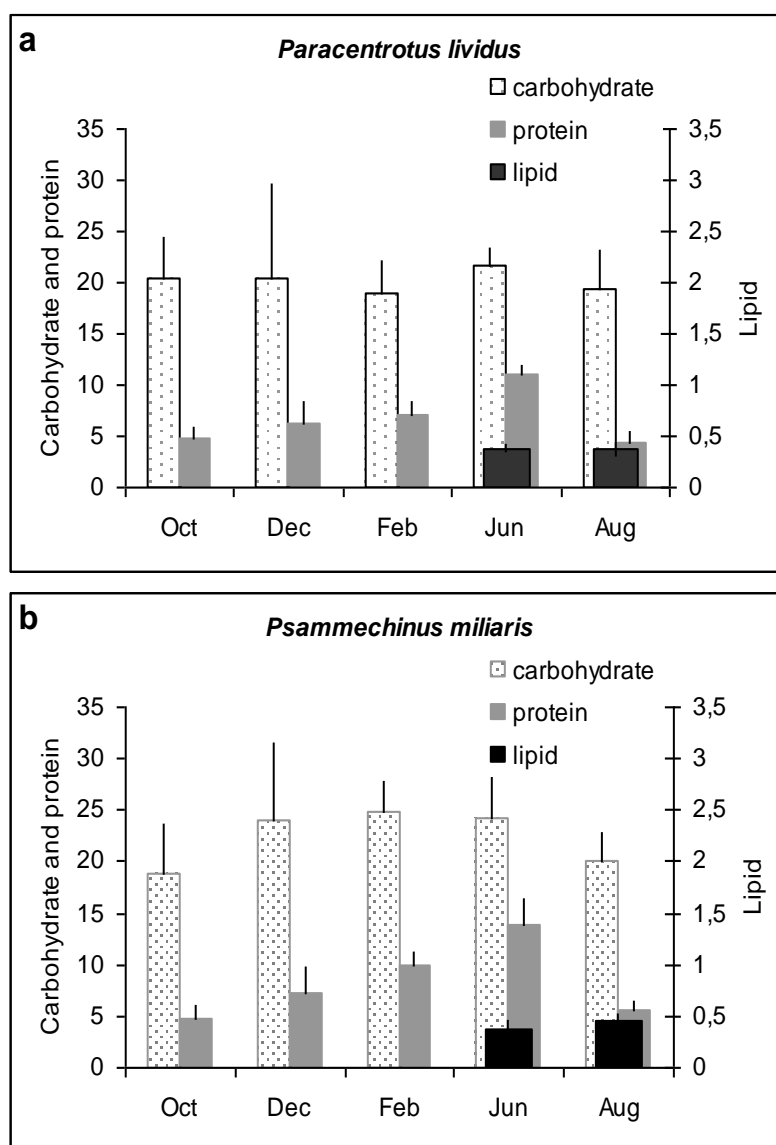


Fig. 7 Seasonal changes in the absorption rate of the sea urchins fed *Palmaria palmata* in term of proteins, carbohydrates and lipids (in mg DW d<sup>-1</sup> urchin<sup>-1</sup>) (+SD). a : *Paracentrotus lividus*; b : *Psammechinus miliaris*.

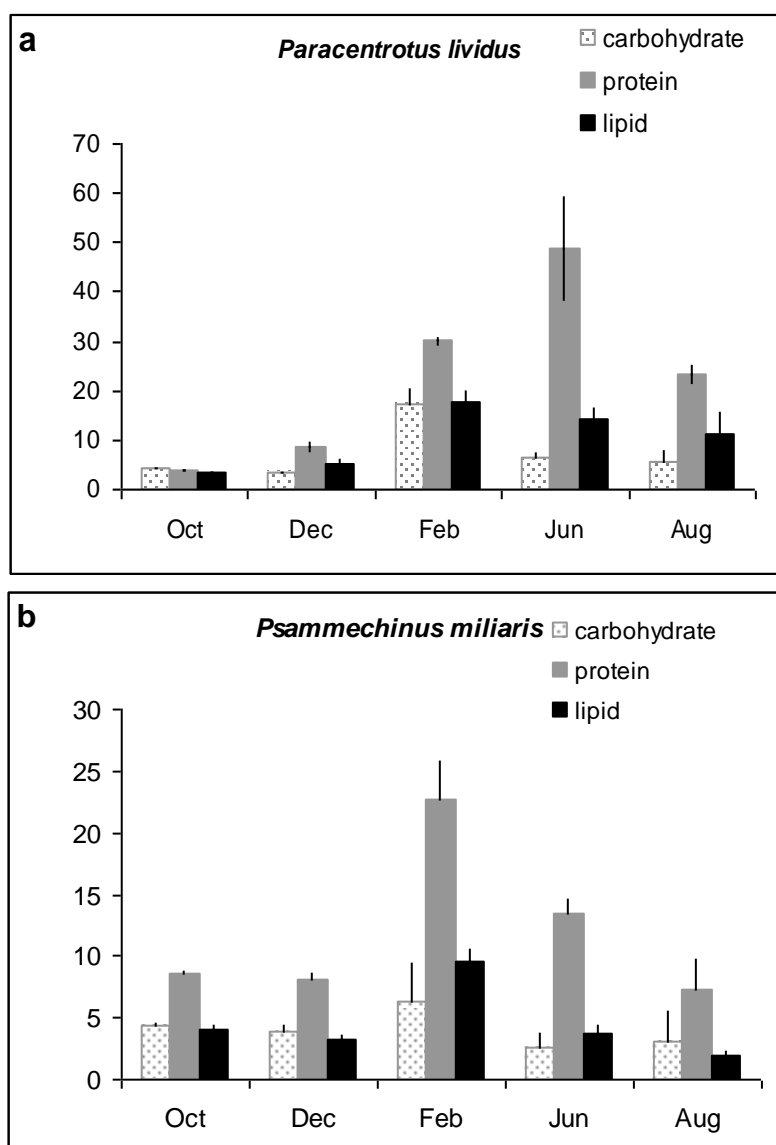


Fig. 8 Seasonal changes in the estimated composition of the gonad of the sea urchins fed *Palmaria palmata* (in mg DW) (+SD). a : *Paracentrotus lividus*; b : *Psammechinus miliaris*.

## REFERENCES

740 **References**

741

742 Akiyama, T., Unuma, T., Yamamoto, T., 2001. Optimum protein level in a  
743 purified diet for young red sea urchin Pseudocentrotus depressus. Fish. Sci.  
744 67, 361-363.

745 Allain, J.Y., 1975. Structure des populations de Paracentrotus lividus  
746 (Lamarck) (Echinodermata, Echinoidea) soumises à la pêche sur les côtes  
747 Nord de Bretagne. Rev. Trav. Inst. Pêches Marit. 39, 171-212.

748 Basuyaux, O., Blin, J.L., 1998. Use of maize as a food source for sea urchin  
749 in a recirculating rearing system. Aquat. Int. 6, 233-247.

750 Bligh, E.G., Dyer, W.J., 1959. A rapid method of total lipid extraction and  
751 purification. Can. J. Biochem. Physiol. 37, 911-917.

752 Boudouresque, C.F., Verlaque, M., 2001. Ecology of Paracentrotus lividus.  
753 In: Lawrence, J.M. (Ed.), Edible Sea Urchins: biology and ecology. Elsevier  
754 Science Publishers B.V., Amsterdam, pp. 177-216.

755 Byrne, M., 1990. Annual reproductive cycles of the commercial sea urchin  
756 Paracentrotus lividus from an exposed intertidal and a sheltered subtidal  
757 habitat on the west coast of Ireland. Mar. Biol. 104, 275-289.

758

759 Chapman, A.R.O., Craige, J.S., 1977. Seasonal growth in Laminaria  
760 longicruris: relations with dissolved inorganic nutrients and internal reserves  
761 of N. Mar. Biol. 40, 197-205.

762

- 763 De Ridder, C., Lawrence, J.M., 1982. Food and feeding mechanisms:  
 764 Echinoidea. In: Jangoux, M., Lawrence, J.M., (Eds), Echinoderm Nutrition.  
 765 A.A. Balkema Publishers, Rotterdam, pp.57-115.  
 766
- 767 Dominique, F., 1973. Contribution à l'étude du cycle annuel de  
 768 reproduction de deux espèces d'échinoides (Echinodermata) des côtes de  
 769 Bretagne. B. SC. Thesis, Université Libre de Bruxelles, Belgique  
 770
- 771 Dubois, M., Gilles, K.A., Hamilton, J.K, Reber, P.A., Smith, F., 1956.  
 772 Colorimetric method for determination of sugar and related substance. Anal.  
 773 Chem. 28, 350-356.  
 774
- 775 Fenaux, L., Malara, G., Cellario, C., Charra, R., Palazzoli, T., 1977.  
 776 Evolution des constituants biochimiques des principaux compartiments de  
 777 l'oursin Arbacia lixula (L.) au cours d'un cycle sexuel et effets d'un jeûne de  
 778 courte durée au cours de la maturation sexuelle. J. Exp. Mar. Biol. Ecol. 28,  
 779 17-30.  
 780
- 781 Fernandez, C., 1996. Croissance et nutrition de Paracentrotus lividus dans le  
 782 cadre d'un projet aquacole avec alimentation artificielle. Thèse de doctorat  
 783 de l'Université de Corse, France.  
 784
- 785 Fernandez, C., 1998. Seasonal changes in the biochemical composition of  
 786 the edible sea urchin Paracentrotus lividus (Echinodermata : Echinoidea) in  
 787 a lagoonal environment. Mar. Ecol. 1998; 19(1), 1-11.

788

789 Fleurence, J., 1999. Seaweed proteins: biochemical , nutritional aspects and  
790 potential uses. Trends Food Sci. Technol. 10, 25-28.

791

792 Frantzis, A., Grémare, A., 1992. Ingestion, absorption, and growth rates of  
793 Paracentrotus lividus (Echinodermata : Echinoidea) fed different  
794 macrophytes. Mar. Ecol. Prog. Ser. 95, 169-183.

795

796 Fuji, A., 1967. Ecological studies on the growth and food consumption of  
797 Japanese common littoral sea urchin, Strongylocentrotus intermedius. Mem.  
798 Fac. Fish., Hokkaido Univ. 15, 83-160.

799

800 Galland-Irmouli, A-V., Fleurence, J., Lamghari, R., Luçon, M., Rouxel, C.,  
801 Barbaroux, O., Bronowicki, J.P., Villaume, C., Guéant, J.L., 1999.  
802 Nutritional value of proteins from edible seaweed Palmaria palmata (Dulse).  
803 J. Nutr. Biochem. 10, 353-359.

804

805 Guillou, M., Grall, J., Connan, S., 2002. Can low sea urchin densities  
806 control macro-epiphytic biomass in a north-east Atlantic maerl bed  
807 ecosystem (Bay of Brest, France)? J. Mar. Biol. Ass. U. K. 82, 867-876.

808

809 Hagen Rødde, R.S., Vårum, K.M., Larsen, B.A., Myklestad, S.M., 2004.  
810 Seasonal and geographical variation in the chemical composition of the red  
811 alga Palmaria palmata (L.) Kuntze. Bot. Mar. 47, 125-133.

812

- 813 Indergaard, M., Minsaas, J., 1991. Animal and human nutrition. In Guiry,  
814 M., Blunden, G. (Eds.), Seaweed resources in Europe: uses and potential. J.  
815 Wiley and Sons Publishers, New-York, pp. 21-64.
- 816
- 817 Kelly, M.S., 2000. The reproductive cycle of the sea urchin Psammechinus  
818 miliaris (Echinodermata : Echinoidea) in a Scottish sea loch. J. Mar. Biol.  
819 Ass. U. K. 80, 909-919.
- 820
- 821 Kelly, M.S., 2001. Environmental parameters controlling gametogenesis in  
822 the echinoid Psammechinus miliaris. J. Exp. Mar. Biol. Ecol. 12, 45-64.
- 823
- 824 Kelly, M.S., Cook, E.J., 2001. The ecology of Psammechinus miliaris. In:  
825 Lawrence, J.M. (Ed.), Edible Sea Urchins: biology and ecology. Elsevier  
826 Science Publishers B.V., Amsterdam, pp. 217-224.
- 827
- 828 Lahaye, M., 1991. Marine algae as source of fibres: determination of soluble  
829 and insoluble dietary fibers contents in some "sea vegetables". J. Sci. Food  
830 Agric. 54, 587-594.
- 831
- 832 Lares, M.T., McClintock, J.B., 1991. The effects of temperature on the  
833 survival, organismal activity, nutrition, growth, and reproduction of the  
834 carnivorous, tropical sea urchin Eucidaris tribuloides. Mar. Behav. Physiol.  
835 19(2), 75-96.
- 836

- 837 Larson, B.R., Vadas, R.L., Keser, M., 1980. Feeding and nutritional ecology  
838 of the sea urchin Strongylocentrotus droebachiensis in Maine, USA. Mar.  
839 Biol. 59, 49-62.
- 840
- 841 Le Gall, L., 2002. Etudes biologiques , biochimiques et cellulaires de  
842 Palmaria palmata (Rhodophyta); applications biotechnologiques à  
843 l'aquaculture. Thèse de doctorat de l'Université de Caen, France.
- 844
- 845 Le Gall, P., 1989. L'échinoculture. Technique et documentation. Lavoisier,  
846 Paris (France). pp.467-491.
- 847
- 848 Le Gall, P., Bucaille, D., Dutot, P., 1989. Résistance aux variations de  
849 salinité chez Paracentrotus et Psammechinus. Vie mar. HS10, 83-84.
- 850
- 851 Le Gall, P., Bucaille, D., Grassin, J.B., 1990. Influence de la température sur  
852 la croissance de deux oursins comestibles, Paracentrotus lividus et  
853 Psammechinus miliaris. In: De Ridder, C, Dubois, P., Lahaye, M.C.,  
854 Jangoux, M. (eds). Echinoderm Research. Balkema publishers, Rotterdam,  
855 pp183-188.
- 856
- 857 Lemire, M., Himmelman, J-H., 1996. Relation of food preference to fitness  
858 for the green sea urchin, Strongylocentrotus droebachiensis. Mar. Biol. 127  
859 (1), 73-78.
- 860



- 861   Lowe, E.F., Lawrence, J.M., 1976. Absorption efficiencies of Lytechinus  
862   variegatus (Lamarck) (Echinodermata: Echinoidea) for selected marine  
863   plants. J. Exp. Mar. Biol. Ecol. 21, 223-234.  
864
- 865   Lowry, O.H., Rosebrough, J.N, Farr, A.L., Randall, R.J., 1951. Protein  
866   measurement with folin reagent. J. Biol. Chem. 193, 265-275.  
867
- 868   Martinez, B., Rico, J.M., 2002. Seasonal variation of P content and major N  
869   pools in Palmaria palmata (Rhodophyta). J. Physiol. 58, 1082-1089.  
870
- 871   Monteiro-Torreiro, M.F., Garcia-Martinez, P., 2003. Seasonal changes in  
872   the biochemical composition of body components of the sea urchin,  
873   Paracentrotus lividus, in Lorbé (Galicia, north-western Spain). J. Mar. Biol.  
874   Ass. U.K. 83, 575-581.  
875
- 876   Morgan, K.C., Wright, J.L.C., Simpson, F.J., 1980. Review of chemical  
877   constituents of the red alga Palmaria palmata (Dulse). Econ. Bot. 34(1), 27-  
878   50.  
879
- 880   Morgan, K.C., Simpson, F.J., 1981. Cultivation of Palmaria palmata. Effect  
881   of light intensity and nitrate supply on growth and chemical composition.  
882   Bot. Mar. 24, 547-552.  
883

- 884 Otero-Villanueva, M., Kelly, M.S., Burnell, G., 2004. How diet influences  
 885 energy partitioning in the regular echinoid Psammechinus miliaris;  
 886 constructing an energy budget. J. Exp. Mar. Biol. Ecol. 304, 159-181.  
 887
- 888 Pearce, C.M., Daggett, T.L., Robinson, S.M.C., 2003. Effects of starch type,  
 889 macroalgal meal source, and  $\beta$ -carotene on gonad yield and quality of the  
 890 green sea urchin, Strongylocentrotus droebachiensis (Müller), fed prepared  
 891 diets. J. Shellfish Res. 17, 1591-1595.  
 892
- 893 Rouxel, C., Bonnabeze, E., Daniel, A., Jerome, M., Etienne, M., Fleurence,  
 894 J., 2001. Identification by SDS PAGE of green seaweeds (Ulva and  
 895 Enteromorpha) used in the food industry. J. Appl. Ecol. 13 (3), 215-219.  
 896
- 897 Sánchez-Machado, D.I., López-Cervantes, J., López-Hernández, J., Paseiro-  
 898 Losada, P., 2004. Fatty acids, total lipid, protein and ash contents of  
 899 processes edible seaweeds. Food Chem. 85, 439-444.  
 900
- 901 Southward, A., Southward, E., 1975. Endangered urchins. New Sci.  
 902 66(944), 70-72.  
 903
- 904 Spirlet, C., Grosjean, P., Jangoux, M., 1998. Reproductive cycle of the  
 905 echinoid Paracentrotus lividus: analysis by means of maturity index. Invert.  
 906 Reprod. Dev. 34, 69-81.  
 907

- 908 Spirlet, C., Grosjean, P., Jangoux, M., 2000. Optimization of gonad growth  
909 by manipulation of temperature and photoperiod in cultivated sea urchins,  
910 Paracentrotus lividus (Lamarck) (Echinodermata). Aquaculture 185, 85-99.  
911
- 912 Strickland, J., Parsons, T., 1972. A practical handbook of seawater analysis.  
913 Fish. Res. Bd. Can. Bull.167, pp.1-310.  
914
- 915 Vadas, R.L., Beal, B., Dowling, T., Fegley, J.C., 2000. Experimental field  
916 tests of natural diets on gonad index and quality in the green sea urchin,  
917 Strongylocentrotus droebachiensis : a case for rapid summer production in  
918 post-spawned animals. Aquaculture 182, 115-135.  
919
- 920 Watts, S.A., Boettger, A., McClintock J.B., Lawrence, J.M., 1998. Gonad  
921 production in the sea urchin Lytechinus variegatus (Lamarck) fed prepared  
922 diets. J. Shellfish Res. 15(5), 1591-1595.  
923  
924